AUG 1 7 2000

510(k) SUMMARY OF SAFETY AND EFFECTIVENESS

K000605

Manufacturing Site:	Stackhouse Incorporated 1100 Bird Center Drive Palm Springs, CA 92262
Contact:	Tom Gutierrez (760) 778-7255 (phone) (760) 778-7274 (fax)
Summary Date	February 15, 2000
Device Trade Name:	Stackhouse Lens Hood Model SA-700/F
Device Common Name	The Stackhouse Lens Hood Model SA-700/F is Stackhouse Incorporated ventilator common name is "Non-sterile surgical gown"
Device Classification Name:	Classified under 79 FYA "Surgical gowns". • 878.4040 Surgical Apparel, 79 FYA
Establishment Registration Number	2020276
Device Class:	Class II
Classification Panel:	Infection Control Device Branch
Predicate Device:	The predicate device is the Stryker Instruments Steri-Shield Surgical Helmet System Hood: MB20 Hood (reference: K944393)
Device Description:	The Stackhouse Lens Hood Model SA-700/Fis non-sterile Surgical Lens and Hood garment. A more complete description of this device and specifications are provided in Section 5.
Intended Use:	The Stackhouse Lens Hood Model SA-700/Fis to be worn by operating room personnel during surgical procedures to protect both the patient and the operating room personnel from the transfer of microorganisms, body fluids and particulate material.
Substantial Equivalence	The Stackhouse Lens Hood Model SA-700/Fis substantially equivalent to the Stryker Steri-Shield Surgical Apparel: Model: MB20 Hood in that: - The intended use is the same - The performance attributes are similar
Summary of Testing and Validation:	The material used in the Stackhouse Lens Hood Model SA-700/Fwere tested in accordance to industry recognized test standards and was validated for the intended use.

DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

AUG 17 2000

Mr. Tom Gutierrez
Regulatory Compliance Engineer
Stackhouse, Incorporated
1100 Bird Center Drive
Palm Springs, California 92262

Re: K000605
Trade Name: Stackhouse Lens Hood Model SA-700/F
Regulatory Class: II
Product Code: FYA
Dated: July 7, 2000
Received: July 7, 2000

Dear Mr. Gutierrez:

We have reviewed your Section 510(k) notification of intent to market the device referenced above and we have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

If your device is classified (see above) into either class II (Special Controls) or class III (Premarket Approval), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 895. A substantially equivalent determination assumes compliance with the Current Good Manufacturing Practice requirements, as set forth in the Quality System Regulation (QS) for Medical Devices: General regulation (21 CFR Part 820) and that, through periodic QS inspections, the Food and Drug Administration (FDA) will verify such assumptions. Failure to comply with the GMP regulation may result in regulatory action. In addition, FDA may publish further announcements concerning your device in the Federal Register. Please note: this response to your premarket notification submission does not affect any obligation you might have under sections 531 through 542 of the Act for devices under the Electronic Product Radiation Control provisions, or other Federal laws or regulations.

This letter will allow you to begin marketing your device as described in your 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801 and additionally 809.10 for in vitro diagnostic devices), please contact the Office of Compliance at (301) 594-4692. Additionally, for questions on the promotion and advertising of your device, please contact the Office of Compliance at (301) 594-4639. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR 807.97). Other general information on your responsibilities under the Act may be obtained from the Division of Small Manufacturers Assistance at its toll-free number (800) 638-2041 or (301) 443-6597 or at its internet address "http://www.fda.gov/cdrh/dsma/dsmamain.html".

Sincerely yours,

Timothy A. Ulatowski
Director
Division of Dental, Infection Control
and General Hospital Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

510 (k) Number (if known): K000605

Device Name: Stackhouse Lens Hood Model SA-700/F

Indication For Use:

The Stackhouse Lens Hood Model SA-700/Fis intended to be worn by operating room personnel during surgical procedures to protect both the patient and the operating room personnel from the transfer of microorganisms, body fluids and particulate material.

(PLEASE DO NOT WRITE BELOW THIS LINE – CONTINUE ON ANOTHER PAGE IF NEEDED)

Concurrence of CDRH, Office of Device Evaluation (ODE)

(Division Sign-Off)
Division of Dental, Infection Control,
and General Hospital Devices
510(k) Number K000605

Prescription Use _____ OR Over-The-Counter Use X
(Per 21 cfr 801.109)

(Optional Format 1-2-96)